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June 19, 2006

Books

Editor, *Disease and Mortality in Sub-Saharan Africa, Second Edition*, Jamison DT, Feachem RG, Makgoba MW, Baingana FK, Bos ER, Hofman KJ and Rogo KO. The World Bank, Washington, D.C., 2006.

Chapters

Leppert M and Hofman K. Fetal Alcohol Syndrome, in *Developmental Disabilities in Infancy and Childhood. Volume II: The Spectrum of Developmental Disabilities*, Chapter 19 pp 281-288. Second Edition. Eds: Capute AJ and Accardo PJ. Paul H. Brookes, Baltimore, MD, 1996.

Hofman K, Anand N, and Keusch G. Bridging the Health Genomics Divide: A Case for Building Research Capacity in the South, in *Genomics, Health and Society Emerging Issues for Public Policy*, pp 143-154. Eds: Knoppers BM and Scriver C. Policy Research Initiative, Canada, 2003.

Durkin M, Schneider H, Pathania V, Nelson K, Solarsh G, Bellows N, Scheffler R, Hofman K. Chapter 49: Learning and Developmental Disabilities, in *Disease Control Priorities in Developing Countries, Second Edition*. The World Bank and Oxford University Press, New York, 2006.

Solarsh G and Hofman K. Chapter 10: Developmental Disabilities, in *Disease and Mortality in Sub-Saharan Africa, Second Edition*. The World Bank, Washington, D.C., 2006.

Monographs (Prepared for Centers for Disease Control, Atlanta, GA)

Efficacy of screening for developmental disabilities

- Down Syndrome
- Neurofibromatosis Type 1

Proceedings (Editor)

Workshop on Screening for Developmental Disabilities in Preschool in South Africa (1997)

Journal Publications

1. Hofman KJ, Milne FJ and Schmidt C: Acne, hypervitaminosis A and hypercalcemia. A case report. *S Afr Med J* 54: 579-580, 1978.
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8. Hofman KJ, Bernhardt BA and Pyeritz RE: Marfan syndrome: Neuropsychological aspects. *Am J Med Genet* 31: 331-338, 1988.
9. Hofman KJ, Antonarakis SE, Missiou-Tsangaraki S, Boehm CD and Valle D: Phenylketonuria in the Greek Population: Haplotype analysis of the phenylalanine hydroxylase gene and identification of a PKU mutation. *Mol Biol Med* 6: 245-250, 1989.
10. Hayflick S, Hofman K, Tunnessen W, Leventhal B and Dugeon D: Neurofibromatosis I: Recognition and Management of Associated Neuroblastoma. *Pediatric Dermatology* 7: 293-295, 1990.
11. Hofman KJ, Steele G, Kazazian HH, and Valle D: Phenylketonuria in US Blacks: Molecular Analysis of the Phenylalanine Hydroxylase Gene. *Am J Hum Genet* 48: 791-798, 1991.
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36. Hofman K, Primack A, Keusch G, and Hrynkow S. Addressing the Growing Burden of Trauma and Injury in Low- and Middle-Income Countries. *Am J Public Health*; 95 (1): 13-17. 2005.
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